CERTIFICATE OF FACSIMILE TRANSMISSION

I hereby certify that this correspondence is being transmitted to the U.S. Patent & Trademark Office

in accordance, with 37 CFR § 1.6(d) on the date indicated.

n__/

ay 27204

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Inventor: M.K. Carpenter et al.

Filing Date: June 21, 2001

Serial No: 09/888,309

Docket: 090/002

Title: DOPAMINERGIC NEURONS OBTAINED

FROM HUMAN EMBRYONIC STEM CELLS

Art Unit: 1632

Examiner: Anne Marie Falk, Ph.D.

DECLARATION UNDER 37 CFR § 1.132 R. SCOTT THIES, Ph.D.

Commissioner for Patents Alexandria, VA 22313-1450

Dear Sir:

I, SCOTT THIES, do hereby declare as follows:

1. I am a cell biologist who has been working in the field of regenerative medicine since 1989. Since 1997, I have been developing procedures to expand and characterize human neural cells. I joined Geron Corporation in March of 2001, and am currently Associate Director of Neurobiology. My project entails the expansion and characterization of neural progenitors from human embryonic stem cells for treatment of conditions such as Parkinson's Disease and Spinal Cord Injury.

A copy of my curriculum vitae accompanies this Declaration.

PATENT 09/888,309 Docket 090/002

- 2. I have reviewed the patent application indicated above, along with the pending claims. The application describes methods and reagents for generating neural cells from pluripotent stem (hES) cells. Example 5 provides data from experiments in which human embryonic stem cells were differentiated into populations that were highly enriched for cells bearing markers of the neural lineage, such as β -tubulin, MAP-2, and tyrosine hydroxylase (Table 5, Figure 4).
- 3. I understand the Examiner has questioned how the combination of undifferentiated hES cells and hES derived neural cells is described in the disclosure, and how these cell populations can be used together.
- 4. The patent application describes reagents, techniques, and strategy for preparing and characterizing populations of specialized differentiated cells. This is referred to as a system, both in the Summary (page 4), and at the outset of the Detailed Description (page 7). The disclosure elaborates how to use the differentiated cells for a variety of purposes, including therapeutic treatment of neurological disease, and drug screening.
- 5. Someone skilled in the production of mammalian cells will understand from reading the disclosure that the "system" referred to encompasses the starting undifferentiated hES cell population, and the various populations that are obtained using the differentiation paradigms that are described and illustrated.

In particular, someone making neural cells by following some of the exemplified methods would begin with a line of undifferentiated hES cells. This meets the requirement of the first component of the claims. As the cells are guided along the neural differentiation pathway as described, they would acquire markers such as β-tubulin, MAP-2, or tyrosine hydroxylase. This meets the requirement of the "second population" indicated in the claims.

6. Someone practicing the invention in this way would therefore have possession of the undifferentiated hES cells, and possession of the cells bearing the neural markers, as part of the system for generating neural progenitor cells or terminally differentiated cells. The neural lineage cells would be suitable for a number of different uses described in the specification, such as the preparation of pharmaceutical compositions, and for use in drug screening.

PATENT 09/888,309 Docket 090/002

F-371

7. It would not be necessary for the user to have possession of the two cell populations at the same time, since a population of undifferentiated hES cells could be caused to differentiate to neural cells in its entirety.

However, a scientist knowledgeable in this area understands the benefit of retaining some of the hES cells in the undifferentiated state, since the undifferentiated hES cells can be further expanded to act as a virtually unlimited reservoir for producing the differentiated cells. The use of the system in this way is provided in the Example section of this patent application. To this day, Geron Corporation continues to maintain hES cell lines in both the undifferentiated form, producing neural cells whenever needed in the quantity required.

8. I hereby declare that all statements made in this Declaration of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.

Menlo Park, California

Geron Corporation 230 Constitution Drive Menlo Park, CA 94025 (650) 473-8657 sthies@geron.com

1053 Riesling Drive Pleasanton, CA 94566 (925) 484-0404 sspt@comcast.net

Education

B.S., University of Miami, Biology and Chemistry, *cum laude*, 1979 Ph.D., Duke University, Physiology, 1983

Honors and Awards

Honors Program, University of Mlami, 1975-1979
Privileged Studies Program, University of Miami, 1976-1979
Alpha Lambda Delta, National Freshman Honors Society, 1976
Beta Beta Beta, National Biology Honors Society, 1977
Phi Kappa Phi, National Honors Society, 1978
Omicron Delta Kappa, National Honors Society, 1978
National Research Service Award, 1979-1983
Sigma Xi Grant-in-Aid of Research, 1980
Biomedical Research Support Grant, 1981
National Research Service Award, 1984-1986
Biomedical Research Support Grant, 1986
National Research Service Award, 1987-1989

Associations

Endocrine Society, Member, 1990-present American Society for Cell Biology, Member, 1991-present Society for Neuroscience, Member, 2000-present International Society for Stem Cell Research, 2004

Teaching Experience

Conference Instructor/Medical Physiology, Duke University, 1980

Professional Experience

Laboratory Technician, Department of Biology University of Miami, 1977-1979

Postdoctoral Fellow, Department of Biological Chemistry University of California, Los Angeles, 1984-1987

Postdoctoral Fellow, Division of Endocrinology and Metabolism, Department of Medicine, University of California, San Diego, 1987-1989

Staff Scientist II, Tissue Growth and Repair, Genetics Institute, Inc., 1989-1993

Principal Scientist, Tissue Growth and Repair, Genetics Institute, Inc., 1993-1997

Senior Scientist, Tissue Growth and Repair, Genetics Institute, Inc., 1997-1998

Senior Scientist, Musculoskeletal Sciences, Genetics Institute/Wyeth, 1998-2001

Group Leader, Stem Cell Therapeutics, Geron Corp., 2001-2003

Radiation Safety Officer, Geron Corp., 2002-present

Associate Director, Stem Cell Therapeutics, Geron Corp., 2003-present

Publications

Abstracts

Thies RS and Mandel LJ (1984) Utilization of endogenous substrates in the rabbit cornea. Fed. Proc. 42(3):674.

Thies RS, Molina JM, Henry RR and Freidenberg GR (1988) Insulin-induced phosphorylation of endogenous substrates in rat and human adipocytes. Diabetes 37(Suppl.1):33A.

McClain DA, Maegawa H, Thies RS, Ullrich A and Olefsky JM (1988) Insulin receptors with defective tyrosine kinase inhibit normal receptor function at the level of substrate phosphorylation. FASEB J. 2(5):A1410.

McClain DA, Thies RS, Maegawa H and Ullrich A (1988) Dissociation of insulin's actions revealed by a truncated insulin receptor. Diabetes 37(Suppl.1):8A.

+6504738654

Thies RS, McClain DA, Maegawa H and Olefsky JM (1989) Dissection of insulin and IGF-1 mediated growth and metabolic signaling in cells expressing kinasedefective insulin receptors. 71st Annual Endocrine Soc. Meeting, Program and Abstracts:46.

Rosen V, Thies RS, Kurtzberg L, D'Allessandro J, Wozney JM and Wang E (1990) In vivo and in vitro activities of recombinant human BMP-2. Calcified Tissue Intl. 46(Suppl.2):45.

Thies RS, Bauduy M, McQuaid D, Kurtzberg L, Cordes P, Capparella J, Wozney JM, Wang E and Rosen V (1990) Bone morphogenetic protein alters W-20 stromal cell differentiation in vitro. J. Bone Min. Res. 5(Suppl.2):305.

Rosen V, Thies RS, Kurtzberg L, Cordes P, McQuaid D, Bauduy M, Moutsatsos I, D'Allessandro J, Wozney JM and Wang E (1990) in vivo and in vitro roles of BMP in skeletal formation and repair. J. Čell Biochem. (Suppl.14E):33.

Rosen V, Bauduy M, McQuaid D, Donaldson D, Thies RS and Wozney JM (1991) Expression of osteoblast-like phenotype in mouse embryo limb bud cell lines cultured in BMP-2 and retinoic acid. 73rd Annual Endocrine Soc. Meeting, Program and Abstracts:151.

Thies RS, Song J, Wozney JM and Rosen V (1992) Vitamin D Inhibits BMP-2 induction of osteocalcin in W-20-17 bone marrow stromal cells. 74th Annual Endocrine Soc. Meeting, Program and Abstracts:443.

Yamaji N, Thies RS, Celeste AJ and Wozney JM (1993) The molecular cloning of bone morphogenetic protein receptors. J. Bone Min. Res. 8(Suppl.1):S145.

Thies RS, Song JJ, Yamaji N, Celeste AJ, Wozney JM and Rosen V (1993) Regulation of BMP receptor affinity by phosphorylating agents. Mol. Biol. Cell 5(Suppl.):295a.

Song JJ, Celeste AJ, Rosen V and Thies RS (1994) Bone Morphogenetic Protein-9 (BMP-9) binds to HepG2 cells and stimulates proliferation. Mol. Biol. Cell 5 (Suppl.):384a.

Neuhaus H, Rosen V and Thies RS (1996) T-Alk, a BMP-type II receptor binds with high affinity to BMP-9. Cold Spring Harbor Meeting on "Mouse Molecular Genetics".

Chevaile A, Santos B, Thies S and Gullans SR (1996) BMP-9, a new member of the TGF-β superfamily, is expressed in the kidney and inhibits proliferation of mesangial and MDCK cells. Int. J. Am. Soc. Nephrol. 7:1654.

Ploemacher RE, Engels LJA, Mayen AEM, Thies S and Neben S (1997) Bone morphogenetic protein-9 is a potent synergistic factor for murine hemopoietic progenitor cell generation and colony formation in serum-free cultures. Blood 90 (Suppl.1):2116.

Chevaile A, Ziai F, Mackenzie HS, Brenner BM, Thies S and Gullans SR (1997) Renal protective effect of bone morphogenetic protein-2 (BMP-2) in a rat model of ischemia-reperfusion injury. Int. J. Am. Soc. Nephrol. 8:2724.

Lopez-Coviella I, Berse B, Thies RS and Blusztajn JK (1998) Bone morphogenetic protein-9 modulates acetylcholine content in the SN56 murine septal cell line. Soc. Neuroscience Abstracts 24:1335.

Neuhaus H, Rosen V and Thies RS (1998) BMP-9 and BMP-10, two novel factors involved in heart and liver organogenesis. Soc. Dev. Biology.

Neuhaus H, Rosen V and Thies RS (1999) Mice deficient for BMP-10 are nonviable due to a defect in heart development. German Soc. Dev. Biology.

Miller AF, Harvey SAK, Thies RS and Olson MS (1999) Bone morphogenetic protein-9 expression and binding in adult rat liver. FASEB J. 13:A684.

Lopez-Coviella I, Berse B, Thies RS and Blusztajn JK (1999) Bone morphogenetic proteins (BMPs) as cholinergic differentiating factors for basal forebrain neurons. Soc. Neuroscience Abstracts 25:517.

Krauss R, Fitzgerald M and Thies RS (2000) BMPs can alter the fate of cortical neuronal precursors. Gordon Conference on "Neuronal Development".

Thies RS (2001) Lineage Specific Differentiation of human embryonic stem cells. ARNMD Conference on "Stem Cells for a New Clinical Neuroscience".

Carpenter MK, Inokuma MS, Denham J, Thies RS, Zaremba A and Rao M (2001) Neural specific differentiation of human embryonic stem cells. Dev. Brain Res. 132 (1,2):A20.

Thies RS (2003) Stem cells, cloning and cellular life span. Gordon Conference on "Biology of Aging".

Carpenter MK, Rosler E, Xu C, Priest C and Thies RS (2003) Human embryonic stem cells: characterization and differentiation. Keystone Symposium on "From Stem Cells to Therapy".

Thies RS, Denham J, Frolkis M, Zhang YW, Murphy B, Zaremba A, Inokuma M, Priest C, Haque N, Polonskaya Y, Lau G, Bronstein A and Carpenter MK (2003) Deriving neural progenitor cells and dopaminergic neurons from human ES cells. Keystone Symposium on "From Stem Cells to Therapy".

Priest CA, Carpenter MK, Denham J, Polonskaya Y and Thies RS (2003) Transplantation of dopaminergic neurons derived from human ES cells into the 6hydroxydopamine rat model of Parkinson's disease. Soc. Neuroscience Abstracts 91:300.2.

Thies RS (2004) Sources of cells for stem cell therapy. University of Rostock Spring School on "Use of Stem Cells in Neurodegenerative Diseases – Basics and Applications"

Li Y, Denham J, Thies S, Brunette E, Majumdar A, Fortin J, Powell SE, Priest C, Keirstead H, Lebkowski J and Mandalam RK (2004) Scalable production of glial progenitor cells from human embryonic stern cells grown in defined animal component-free culture. Annual meeting of Am. Inst. Chem. Eng.

<u>Papers</u>

Thies RS and Mandel LJ (1985) Role of glucose in corneal metabolism. Am. J. Physiol. 249:C409-C416.

Maegawa H, Olefsky JM, Thies, RS, Boyd D, Ullrich A and McClain DA (1988) Insulin receptors with defective tyrosine kinase inhibit normal receptor function at the level of substrate phosphorylation. J. Biol. Chem. 263:12629-12637.

Thies RS, Ullrich A and McClain DA (1989) Augmented mitogenesis and impaired metabolic signaling mediated by a truncated insulin receptor. J. Biol. Chem. 264:12820-12825.

Thies RS, Molina JM, Ciaraldi TP, Freidenberg GR and Olefsky JM (1990) Insulin receptor autophosphorylation and endogenous substrate phosphorylation in human adipocytes from control, obese and non-insulin dependent diabetic subjects. Diabetes 39:250-259.

McClain DA, Maegawa H, Thies RS and Olefsky JM (1990) Dissection of the growth versus metabolic effects of insulin and insulin-like growth factor-l in transfected cells expressing kinase-defective human insulin receptors. J. Biol. Chem. 265:1678-1682.

Thies RS, Webster NJ and McClain DA (1990) A domain of the insulin receptor required for endocytosis in rat fibroblasts. J. Biol. Chem. 265:10132-10137.

+6504738654

Lima FB, Thies RS and Garvey WT (1991) Glucose and insulin regulate insulin sensitivity in primary cultured adipocytes without affecting insulin receptor kinase activity. Endocrinology 128:2415-2426.

Tucker BJ, Anderson CM, Thies RS, Collins RC and Blantz RC (1992) Glomerular hemodynamic alterations during acute hyperinsulinemia in normal and diabetic rats. Kidney Int. 42:1160-1168.

Thies RS, Bauduy M, Ashton BA, Kurtzberg L, Wozney JM and Rosen V (1992) Recombinant human bone morphogenetic protein-2 induces osteoblastic differentiation in W-20-17 stromal cells. Endocrinology 130:1318-1324.

Rosen V and Thies RS (1992) The BMP proteins in bone formation and repair. Trends Genet. 8:97-102.

Suzuki A, Thies RS, Yamaji N, Song JJ, Wozney JM, Murakami K and Ueno N (1994) A truncated BMP receptor affects dorsal-ventral patterning in early Xenopus embryo. Proc. Natl. Acad. Sci. U.S.A. 91:10255-10259.

Rosen V, Nove J, Song JJ, Thies RS, Cox K and Wozney JM (1994) Responsiveness of clonal limb bud cell lines to BMP-2 reveals a sequential relationship between cartilage and bone cell phenotypes. J. Bone Min. Res. 9:1759-1768.

Graff JM, Thies RS, Song JJ, Celeste AJ and Melton DA (1994) Studies with a Xenopus BMP receptor suggest that ventral mesoderm-inducing signals override dorsal signals in vivo. Cell 79:169-179.

Yamaji N, Celeste AJ, Thies RS, Song JJ, Bernier SM, Goltzman D, Lyons KM, Nove J, Rosen V and Wozney JM (1994) A mammalian serine/threonine kinase receptor specifically binds BMP-2 and BMP-4. Biochem. Biophys. Res. Comm. 205:1944-1951.

Song JJ, Celeste AJ, Kong FM, Jirtle RL, Rosen V and Thies RS (1995) Bone morphogenetic protein-9 binds to liver cells and stimulates proliferation. Endocrinology 136:4293-4297.

Rosen V, Thies RS and Lyons KM (1996) Signaling pathways in skeletal formation: A role for BMP receptors. Ann. N.Y. Acad. Sci. 785:59-69.

Ploemacher RE, Engels LJA, Mayen AEM, Thies RS and Neben S (1999) Bone morphogenetic protein-9 is a potent synergistic factor for a murine hemopoletic progenitor cell generation and colony formation in serum-free conditions. Leukemia 13:428-437.

Neuhaus H, Rosen V and Thies RS (1999) BMP-10 is specifically expressed in the trabeculated part of the myocardial wall. Mech. Dev. 80:180-184.

Miller AF, Harvey SAK, Thies RS and Olson MS (2000) Bone morphogenetic protein-9: An autocrine/paracrine cytokine in the liver. J. Biol. Chem. 275:17937-17945.

Lopez-Coviella I, Berse B, Krauss R, Thies RS and Blusztajn JK (2000) Induction and maintenance of the neuronal cholinergic phenotype in the central nervous system by BMP-9. Science 289:313-316.

Thies RS, Chen T, Davies MV, Tomkinson KN and Wolfman NM (2001) GDF-8 propeptide binds to GDF-8 and antagonizes biological activity by inhibiting GDF-8 receptor binding. Growth Factors 18:251-259.

Lopez-Coviella I, Berse B, Thies RS and Blusztajn JK (2002) Upregulation of acetylcholine synthesis by bone morphogenetic protein-9 in a murine septal cell line. J. Physiol. Paris 96:53-59.

Bhattacharya B, Miura T, Brandenderg R, Mejido J, Luo Y, Yang AX, Joshi BH, Irene G, Thies RS, Amit M, Lyons I, Condie BG, Iskovitz-Elder J, Rao MS and Puri RK (2004) Gene expression in human embryonic stem cell lines: Unique molecular signature. Blood 103(8):2956-2964.

Ginis I, Luo Y, Miura T, Thies S, Brandenberger R, Gerecht-Nir S, Ami M, Hoke A, Carpenter MK, Itskovitz-Eldor J, Rao MS (2004) Differences between human and mouse embryonic stem cells. Dev. Biol. 269: 360-380.

Brandenberger R, Khrebtukova I, Thies RS, Miura T, Puri R, Vasicek T, Lebkowski J and Rao M (2004) MPSS profiling of human embryonic stem cells. Submitted

Chapters and Books

Rosen V, Capparella J, McQuaid D, Cox K, Thies RS, Song J and Wozney JM (1993) Development of Clonal Cell Lines Derived From 13 DPC Mouse Limb Buds as a System to Study the Effects of Recombinant Human BMP-2 on Limb Bud Cell Differentiation. In Limb Development and Regeneration; Fallon JF, Goetinck PF, Kelley RO and Stocum DL, Ed.; Wiley-Liss: New York, 305-315.

Rosen V and Thies RS (1993) Skeletal Formation and the Bone Morphogenetic Proteins. In Growth Factors in Perinatal Development; Tsang RC, Lemons JA and Balistreri WF, Ed.; Raven Press: New York, 39-58.

Rosen V and Thies RS (1995) The Cellular and Molecular Basis of Bone Formation and Repair; RG Landes Co.: Austin.

<u>Patents</u>

Wozney JM, Celeste AJ, Thies RS and Yamaji N (1994) Activin receptor-like kinase (ALK), belonging to the TGF receptor family and/or to the BMP receptor family. International Patent WO9507982.

Rosen V, Wozney JM, Celeste AJ, Thies RS and Song JJ (1995) BMP-9 compositions. International Patent WO9533830.

Celeste AJ, Wozney JM and Thies RS (1999) Neuronal uses of BMP-11. International Patent WO9924057.

Celeste AJ, Wozney JM and Thies RS (1999) Neuronal uses of BMP-11. International Patent WO9924058.

Rosen V, Wozney JM, Celeste AJ, Thies RS and Song JJ (2000) BMP-9 compositions. United States Patent 6034061.

Thies RS and Song JJ (2000) Bone morphogenetic protein (BMP)-9 compositions and their uses. United States Patent 6034062.

Wozney JM, Celeste AJ, Thies RS and Yamaji N (2001) BMP receptor proteins. United States Patent 6291206.

Rosen V, Wozney JM, Celeste AJ, Thies RS and Song JJ (2001) BMP-9 compositions. United States Patent 6287816.

Blusztajn JK, Lopez-Coviella I, Krauss R and Thies RS (2002) BMP-9 compositions and methods for inducing differentiation of cholinergic neurons. International Patent WO0117547.

Celeste AJ, Wozney JM and Thies RS (2002) Neuronal uses of BMP-11. United States Patent 6340668.

Xu C and Thies RS (2003) Mesenchymal cells and osteoblasts from human embryonic stem cells. International Patent WO03004605.

Carpenter MK, Denham JJ, Inokuma MS and Thies RS (2003) Dopaminergic neurons and proliferation-competent precursor cells for treating Parkinson's disease. International Patent WO03000868.

J. Michael Schiff geron GERON CORPORATION 230 Constitution Drive Menlo Park, CA 94025 Phone: (650) 473-7715

Fax: (650) 473-8654

Last page of this transmission